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(54) Title: METHODS AND MATERIALS RELATING TO DNA BINDING PROTEINS

(57) Abstract

DNA sequences associated with regulation of early stages of cell growth are described. Illustratively provided are human and mouse origin DNA sequence encoding early growth regulatory ("Egr") proteins which include "zinc finger" regions of the type involved in DNA binding. Immunological methods and materials for detection of Egr proteins and hybridization methods and materials for detection and quantification of Egr protein related nucleic acids are also disclosed.

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"METHODS AND MATERIALS RELATING TO DNA BINDING PROTEINS"

CROSS-REFERENCE TO RELATED APPLICATIONS

This is a continuation-in-part application of co-pending U.S. Patent Application Serial No. 179,587, filed April 8, 1988.

BACKGROUND OF THE INVENTION

The present invention relates generally to DNA binding regulatory proteins and more particularly to DNA sequences encoding early growth regulatory proteins possessing histidine-cysteine "zinc finger" DNA binding domains, to the polypeptide products of recombinant expression of these DNA sequences, to peptides and polypeptides whose sequences are based on amino acid sequences deduced from these DNA sequences, to antibodies specific for such proteins and peptides, and to procedures for detection and quantification of such proteins and nucleic acids related thereto.

Among the most significant aspects of mammalian cell physiology yet to be elucidated is the precise manner in which growth factors (e.g., hormones, neurotransmitters and various developmental and differentiation factors) operate to effect the regulation of cell growth. The interaction of certain growth factors with surface receptors of resting cells appears to rapidly induce a cascade of biochemical events thought to result in nuclear activation of specific growth related genes, followed by ordered expression of other genes. Analysis of sequential activation and expression of genes during the transition from a resting state ("G_0") to the initial growing state ("G_1") has been the subject of substantial research. See, gener-

ally, Lau et al., Proc. Nat'l. Acad. Sci. (USA), 84, 1182-1186 (1987). Much of this research has involved analysis of the expression of known genes encoding suspected regulatory proteins (such as the proto-oncogenes, c-fos and c-myc) following mitogen stimulation. An alternative approach has involved attempts to identify genes activated by mitogenic stimuli through differential screening of cDNA libraries prepared from resting cells following exposure to serum and specific growth factors. See, e.g., Lau et al., EMBO Journal, 4, 3145-3151 (1985). See also, Cochran et al., Cell, 33, 939-947 (1983), relating to the cloning of gene sequences apparently regulated by platelet derived growth factor.

Of interest to the background of the invention is the continuously expanding body of knowledge regarding structural components involved in the binding of regulatory proteins to DNA. Illustratively, the so-called receptor proteins are believed to bind to DNA by means of zinc ion stabilized secondary structural fingers premised on folding of continuous amino acid sequences showing high degrees of conservation of cysteines and histidines and hydrophobic residues. See, e.g., Gehring, TIBS, 12, 399-402 (1987). For example, a "zinc finger" domain or motif, present in Xenopus transcription factor IIIA (TF IIIA), as well as the Drosophila Kruppel gene product and various yeast proteins, involves "repeats" of about 30 amino acid residues wherein pairs of cysteine and histidine residues are coordinated around a central zinc ion and are thought to form finger-like structures which make contact with DNA. The histidine-cysteine (or "CC-HH") zinc finger motif, as opposed to a cysteine-cysteine ("CC-CC") motif of steroid receptors, is reducible to a consensus sequence represented as $C-X_{2-4}-C-X_3-F-X_5-L-X_2-H-X_3-H$ wherein C represents cysteine, H represents

histidine, F represents phenylalanine, L represents leucine and X represents any amino acid. [See, Klug et al. TIBS, 12, 464-469 (1987); Blumberg et al., Nature, 328, 443-445 (1987); and Schuh et al., Cell, 47, 1025-1032 (1986).]

Of particular interest to the background of the invention is the recent report of Chowdhury et al., Cell, 48, 771-778 (1987), relating to an asserted "family" of genes encoding proteins having histidine-cysteine finger structures. These genes, designated "mkr1" and "mkr2", appear to be the first such isolated from mammalian tissue and are not correlated to any early growth regulatory events.

There continues to exist a need in the art for information concerning the primary structural conformation of early growth regulatory proteins, especially DNA binding proteins, such as might be provided by knowledge of human and other mammalian DNA sequences encoding the same. Availability of such DNA sequences would make possible the application of recombinant methods to the large scale production of the proteins in procaryotic and eukaryotic host cells, as well as DNA-DNA and DNA-RNA hybridization procedures for the detection, quantification and/or isolation of nucleic acids associated with these and related proteins. Possession of such DNA-binding proteins and/or knowledge of the amino acid sequences of the same would allow, in turn, the development of monoclonal and polyclonal antibodies thereto (including antibodies to protein fragments or synthetic peptides modeled thereon) for use in immunological methods for the detection and quantification of early growth regulatory proteins in fluid and tissue samples as well as for tissue specific delivery of substances such as labels and therapeutic agents to cells expressing the proteins. In addition, DNA probes based on the DNA sequences for these mammalian early growth

regulatory proteins may be of use in detecting gene markers used for the diagnosis of those clinical disorders which are linked to the marker genes.

BRIEF SUMMARY OF THE INVENTION

The present invention provides novel purified and isolated DNA sequences encoding mammalian early growth regulatory ("Egr") proteins which comprise one or more histidine-cysteine zinc finger amino acid sequences putatively providing DNA binding (and hence DNA replication or transcription regulatory) capacity. In presently preferred forms, novel DNA sequences of the invention comprise genomic and cDNA sequences encoding human and mouse early growth regulatory proteins. Alternate DNA forms, such as "manufactured" DNA, prepared by partial or total chemical synthesis from nucleotides, are also within the contemplation of the invention.

Operative association of Egr-encoding DNA sequences provided by the invention with homologous or heterologous species expression control DNA sequences, such as promoters, operators, regulators and the like, allows for in vivo and in vitro transcription to form messenger RNA which, in turn, is susceptible to translation to provide Egr proteins in large quantities. In one presently preferred DNA expression system practiced according to the invention, Egr-encoding DNA is operatively associated with a bacteriophage T3 or T7 RNA promoter DNA sequence allowing for in vitro transcription and translation in a cell free system. Incorporation of novel DNA sequences of the invention into procaryotic and eucaryotic host cells by standard transformation and transfection processes involving suitable viral and circular DNA plasmid vectors is also within the contemplation of the invention and is expected to

provide useful proteins in quantities heretofore unavailable from natural sources. Illustratively, fragments of DNA encoding Egr protein of the invention have been incorporated in plasmid vectors resulting in expression by transformed E.coli hosts of fusion proteins sharing immunological characteristics of Egr protein. Use of mammalian host cells is expected to provide for such post-translational modifications (e.g., truncation, glycosylation, and tyrosine, serine or threonine phosphorylation) as may be needed to confer optimal biological activity on recombinant expression products of the invention.

Also provided by the present invention are novel, presumptively mitogen sensitive, DNA sequences involved in regulation of the transcription of Egr-encoding DNA, which sequences are expected to have utility in the efficient recombinant expression of Egr proteins as well as proteins encoded by other structural genes. In addition, the DNA sequences may be used as probes to detect the presence or absence of gene markers used for the diagnosis of clinical disorders linked to those gene markers.

Novel polypeptide products of the invention include polypeptides having the primary structural conformation (i.e., amino acid sequence) of Egr proteins or fragments thereof, as well as synthetic peptides, an analogs thereof, assembled to be partially or wholly duplicative of amino acid sequences extant in Egr proteins. Proteins, protein fragments, and synthetic peptides of the invention are expected to have therapeutic, diagnostic, and prognostic uses and also to provide the basis for preparation of monoclonal and polyclonal antibodies specifically immunoreactive with Egr proteins, as well as to provide the basis for the production of drugs for use as competitive inhibitors or potentiators of Egr. Preferred protein fragments and

synthetic peptides of the invention include those duplicating regions of Egr proteins which are not involved in DNA binding functions (i.e., regions other than the zinc fingers). Most preferred are peptides which share at least one continuous or discontinuous antigenic epitope with naturally occurring Egr proteins.

Antibodies of the invention preferably bind with high immunospecificity to Egr proteins, fragments, and peptides, preferably recognizing epitopes which are not common to other proteins, especially other DNA binding proteins.

Also provided by the present invention are novel procedures for the detection and/or quantification of Egr proteins and nucleic acids (e.g., DNA and mRNA) specifically associated therewith. Illustratively, antibodies of the invention may be employed in known immunological procedures for quantitative detection of Egr proteins in fluid and tissue samples. Similarly, DNA sequences of the invention (particularly those having limited homology to other DNAs encoding DNA binding proteins) may be suitably labelled and employed for the quantitative detection of mRNA encoding the proteins. Information concerning levels of Egr mRNA may provide valuable insights into growth characteristics of cells.

Among the multiple aspects of the present invention, therefore, is the provision of (a) novel purified and isolated Egr-encoding DNA sequences set out in Figures 1A, 3, and 4 as well as (b) Egr-encoding DNA sequences which hybridize thereto under hybridization conditions of the stringency equal to or greater than the conditions described herein and employed in the initial isolation of DNAs of the invention, and (c) synthetic or partially synthetic DNA sequences encoding the same, or allelic variant, or analog Egr polypeptides

which employ, at least in part, degenerate codons. Correspondingly provided are viral or circular plasmid DNA vectors incorporating such DNA sequences and procaryotic and eucaryotic host cells transformed or transfected with such DNA sequences and vectors as well as novel methods for the recombinant production of Egr proteins through cultured growth of such hosts and isolation from the hosts or their culture media.

Preferred polypeptide products of the invention include those wholly or partially duplicating the deduced sequence of the amino acid residues set out in Figures 1A and 3 (i.e., mouse "Egr-1" and human "EGR2"). Other preferred polypeptides include fusion proteins such as cro-ß-galactosidase/Egr-1 and bovine growth hormone/Egr-1.

Presently preferred antibodies of the invention include those raised against synthetic peptides partially duplicating deduced Egr amino acid sequences of Figures 1A and 3 (e.g., the synthetic peptides H-L-R-Q-K-D-K-K-A-D-K-S-C, the first 12 amino acid residues of which duplicate mouse Egr-1 residues 416-427 with the last cysteine added for coupling to KLH; and C-G-R-K-F-A-R-S-D-E-R-K-R-H-T-K-I duplicating mouse Egr-1 residues 399-415). The antisera against the first peptide is designated VPS10 and comprises a preferred antibody of the invention.

As employed herein, the term "early growth regulatory protein" shall mean and include a mammalian DNA binding protein encoded by DNA whose transcription temporally corresponds to cellular events attending the G_0/G_1 growth phase transition. As employed herein, "histidine-cysteine zinc finger amino acid sequence" shall mean and include the following sequence of amino acids $C-X_{2-4}-C-X_3-F-X_5-L-X_2-H-X_3-H$ wherein C represents cysteine, H represents histidine, F represents phenylalanine, L represents lysine, and X represents an amino acid.

Other aspects and advantages of the present invention will be apparent upon consideration of the following detailed description thereof which includes numerous illustrative examples of the practice of the invention, reference being made to the drawing wherein:

Figure 1A provides a 3086 base nucleotide sequence for a mouse Egr-1 DNA clone as well as a deduced sequence of 533 amino acid residues for the protein; Figure 1B provides a partial restriction map of Egr-1 DNA clones together with information concerning the position of the protein coding sequence and the locus of amino acids providing for histidine-cysteine zinc fingers;

Figure 2 provides an amino acid sequence alignment of the DNA binding domain of mouse Egr-1 in comparison with a zinc finger consensus sequence, with the Drosophila Kruppel sequence and with the "finger 2" sequence of Xenopus TFIIIA protein;

Figure 3 provides a 2820 base nucleotide sequence for a human EGR2 cDNA clone as well as a deduced sequence of 456 amino acids for the protein;

Figure 4 provides a 1200 base nucleotide sequence of a mouse Egr-1 genomic clone, specifically illustrating the 5' non-transcribed regulatory region thereof comprising bases -935 through +1; and

Figure 5 provides a restriction map and organization of the mouse Egr-1 genomic clone mgEgr-1.1 and a comparison to mouse Egr-1 cDNA.

DETAILED DESCRIPTION

The following examples illustrate practice of the invention. Example 1 relates to the preparation and structural analysis of cDNA for mouse Egr-1. Example 2 relates to confirmation of the presence of an Egr DNA sequence on human chromosome 5. Example 3 relates to the *in vitro* transcription and translation of mouse

Egr-1 cDNA. Example 4 relates to production of antibodies according to the invention. Example 5 relates to the isolation and characterization of genomic DNA which encodes mouse Egr-1. Example 6 relates to the isolation and characterization of cDNA encoding human EGR2. Example 7 relates to preparation, in an E. coli host, of a recombinant fusion protein including a portion of the deduced amino acid sequence of mouse Egr-1. Example 8 relates to use of DNA probes of the invention in the quantitative detection of EGR1 mRNA.

These examples are for illustrative purposes only and are not intended in any way to limit the scope of the invention.

EXAMPLE 1

Preparation and Structural Analysis of cDNA for Mouse Egr-1

Isolation of DNA encoding a mammalian early growth regulatory protein including one or more histidine-cysteine zinc finger amino acid sequences was performed substantially according to the procedures described in Sukhatme et al., Oncogene Research, 1, 343-355 (1987), the disclosures of which are specifically incorporated by reference herein.

Balb/c 3T3 cells (clone A31) from the American Type Culture Collection were grown to confluence in Dulbecco's Modified Eagle's medium (DME) supplemented with 10% fetal calf serum (FCS). The cells were rendered quiescent by reduction of the serum concentration to 0.75% for 48 hours. To induce the cells from quiescence into growth phase G_1, the medium was changed to 20% FCS with cycloheximide added to a final concentration of 10 µg/ml.

RNA was extracted from Balb/c 3T3 cells harvested three hours after induction of quiescent cells by

20% FCS and 10 μg/ml cycloheximide. A λgt10 cDNA library was constructed from this mRNA according to the procedures of Huynh et al., DNA Cloning, Vol. 1, 49-78 (Glover, D., ed., IRL Press, 1985). This library was screened differentially with single stranded cDNA prepared from quiescent cells and from cells exposed to serum and cycloheximide for 3 hours. These ^{32}P-labeled cDNA probes were prepared from poly A^+ RNA as described in St. John, et al., Cell, 16, 443-452 (1979), except that 100 μCi of ^{32}P-dCTP (>3000 Ci/mmol), 0.02 mM cold dCTP and 2-5 μg of poly A^+ RNA was used in each reaction. The mean size of the reverse transcribed probes, as assessed by alkaline agarose gel electrophoresis and subsequent autoradiography, was about 700 bases. Replica filter lifts (GeneScreenPlus, NEN-DuPont) were prepared essentially as described by Benton et al., Science, 196, 180-192 (1977), and approximately 3 x 10^6 cpm of ^{32}P-cDNA were used per filter (90 mm diameter). Hybridizations were carried out at 65°C in 1% SDS, 10% dextran sulfate, and 1 M NaCl for a period of 16 hours. The filters were washed twice for twenty minutes each time, first at room temperature in 2 x SSC [Maniatis et al., Molecular Cloning, Cold Spring Harbor Laboratory (New York, 1982)], then at 65°C in 2 x SSC, 1% $NaDodSO_4$ and finally at 65°C in 0.2 x SSC. Autoradiograms were prepared by exposing the blots for 18 hours at -70°C with an intensifying screen.

A total of 10,000 cDNA clones from the Balb/c 3T3 λgt10 library were differentially screened. Seventy-eight clones were found to hybridize preferentially to single-stranded cDNA from fibroblasts stimulated for 3 hours with 20% FCS and cycloheximide as compared to single-stranded cDNA from quiescent cells. Inserts from these clones were cross-hybridized to each other, resulting in the sorting of forty clones into 7 cDNA families one of which was identified as c-fos.

Another cDNA clone, referred to as OC68, contained a 2.2 kb insert and was characterized further. This insert was subcloned into the Eco RI site of pUC13 and probes were generated for Northern blot analysis either from the insert or the corresponding pUC plasmid. Figure 1B illustrates a partial restriction digest map of the OC68 clone ("R" representing restriction sites for RsaI) along with that of a shorter clone, OC19t. Two RsaI digestion fragments, derived from the 5' end of clone OC68 and each comprising approximately 130 base pairs, were labeled and employed to re-screen the above-described λgt10 cDNA library, resulting in the recovery of a 3.1 kb clone, designated OC3.1, shown in figure 1B. This clone was sequenced according to the method of Sanger et al., Proc. Nat'l. Acad. Sci. (USA), 74, 5463 (1977). The 3086 base pair sequence obtained is set forth in Figure 1A along with the deduced sequence of 533 amino acid residues for the protein encoded, designated mouse "Egr-1".

The deduced amino acid sequence shows a single long open reading frame with a stop codon (TAA) at position 1858. The most 5', in-frame, ATG, at position 259, is flanked by sequences that fulfill the Kozak criterion ($\binom{A}{G}$NN(ATG)G) [Kozak, Nuc. Acids Res., 15, 8125-8131 (1987)]. The sequence region upstream of this ATG is highly GC-rich and results in an absence of in-frame stop codons. The 3' untranslated region (UT) contains two "AT" rich regions (nucleotides 2550-2630 and 2930-2970). Similar sequences are found in the 3' UT regions of several lymphokine and proto-oncogene mRNAs, including granulocyte macrophage colony stimulating factor (GM-CSF), interleukin 1, interleukin 2, interleukin 3 (IL-3), α, β, and γ interferons, and c-fos, c-myc, and c-myb [Shaw et al., Cell, 46, 659-667 (1986)]. These sequences may mediate selective mRNA degradation. The presence in the mouse Egr-1 transcript

of such regions is consistent with its short message half-life. Potential polyadenylation signals (AATAAA) are located at nucleotide positions 1865 and 3066, as well as at position 3053 (AATTAA) [Wickens et al., Science, 226, 1045-1051 (1984)].

The deduced amino acid sequence predicts a polypeptide of 533 amino acids with a molecular weight of 56,596. Based on structural considerations, namely a central region containing zinc fingers (described below), the Egr-1 protein can be divided into three domains. The N-terminal portion (amino acid residues 2 to 331) is rich in proline (14.2%) and serine (16%) residues with 7.9% alanines and 7.9% threonines. The C-terminal region (residues 417 to 533) also contains a very high proportion of prolines and serines (15.4 and 26.5%, respectively) and 10.3% alanines and 11.1% threonines. The large number of proline residues leads to a secondary structure that probably lacks α-helices. The central portion of the Egr-1 protein consists of three tandem repeat units of 28-30 amino acids, with the first unit starting at position 332. Each unit conforms almost exactly to the consensus sequence $TGX_{3}^{Y}{}_{F}XCX_{2-4}CX_3FX_5LX_2HX_3H$ (see Figure 2), diagnostic of DNA binding zinc fingers [Berg, Science, 232, 485-486 (1986); Brown et al., Nature, 324, 215 (1986); and Brown et al., FEBS Letters, 186, 271-274 (1985)]. Furthermore, the Egr-1 fingers are connected by "H-C links" ($TGE^{R}_{K}P^{F}_{Y}X$) [Schuh et al., Cell, 47, 1025-1032 (1986)] found in the Xenopus TFIIIA gene (between fingers 1, 2, and 3), in the Drosophila Kruppel gap gene [Rosenberg et al., Nature, 319, 336-339 (1986)], and in genes from mouse and Xenopus that cross-hybridize to the Kruppel (Kr) finger domains: mkr1, mkr2 [Chowdhury et al., Cell, 48, 771-778 (1987)], and Xfin [Altaba et al., EMBO Journal, 6, 3065-3070 (1987)]. The sequence similarity amongst the Egr-1

fingers is 50-70%, whereas the sequence similarity between any of the Egr-1 fingers and those present in TFIIIA, Kruppel, mkr1, mkr2 or Xfin is 35-40%. Outside of the finger domains, it is noteworthy that the Egr-1 and Kr proteins each contain a very high proportion of Pro, Ala, and Ser residues [Schuh et al., Cell, 47, 1025-1032 (1986)]. However, there is no sequence similarity in these regions. Thus, Egr-1 and Kr are not homologous genes nor is Egr-1 related to mkr1, mkr2, Xfin, or TFIIIA. The Kr gene contains thirteen copies of the hexanucleotide (ACAAAA), or its complementary sequence, eight of which are located within 180 bp downstream from the Kr TATA box and five are in the 3' UT region. These sequences may serve as targets for other DNA binding proteins or in Kr gene autoregulation. The Egr-1 cDNA also contains nine copies of the ACAAAA sequence or its complement.

Following the work described above, Milbrandt [Science, 238, 797-799 (1987)], reported the isolation and sequence of a nerve growth factor (NGF) inducible cDNA (NGFI-A) from the rat pheochromocytoma PC12 line. A comparison of the deduced amino acid sequence of NGFI-A to that of mouse Egr-1 of Figure 1A reveals 98% sequence identity. Thus, mouse Egr-1 and rat NGFI-A are homologs. The putative initiation ATG chosen by Milbrandt corresponds to position 343 in the Figure 1A cDNA sequence, and is 84 nucleotides (28 amino acid residues) downstream of the ATG therein designated for translation initiation. Both ATG's have a purine at position -3 and a G at position +1 and the designation represented in Figure 1A of the more 5' ATG as the putative start codon is based on the experience of Kozak, Nuc. Acids Res., 15, 8125-8131 (1987), even though the more 3' ATG is surrounded by the longer Kozak consensus sequence (CCG/ACCATGG). Translation of an *in vitro* generated RNA transcript, described *infra*, selects the more 5' ATG for initiation.

It is noteworthy that a major difference in the deduced sequences of mouse Egr-1 and rat NGFI-A resides in the sequence spanning residues 61-68 of Egr-1 and 33-43 of NGFI-A. The former includes the sequence N-S-S-S-S-T-S-S while the latter includes the sequence N-N-S-S-S-S-S-S-S-S-S, accounting for the 3 residue difference in length of the putative polypeptides which is not accounted for by the difference in designation of the transcript initiation signal.

EXAMPLE 2

Human Chromosome Gene Mapping

To determine the human chromosomal localization of the gene corresponding to mouse Egr-1, the OC3.1 and OC19t cDNA clones were hybridized to a panel of rodent x human somatic cell hybrids. Southern blot analysis of the hybrid panel showed concordance between the presence of Egr-1 sequences and human chromosome 5. In situ hybridization to normal human metaphase chromosomes resulted in specific labeling only of chromosome 5, with the largest cluster of grains at 5q23-31. Specific labeling of these bands was also observed in hybridizations using an Egr-1 probe which does not contain finger sequences.

This localization is interesting in light of the non-random deletions [del(5q)] in human myeloid disorders (acute myelogenous leukemia) (AML), and myelodysplastic syndromes, that involve this chromosomal region. [Le Beau et al., Science, 231, 984-987 (1986); Dewald et al., Blood, 66, 189-197 (1985); and Van den Berghe et al., Cancer Genet. Cytogenet., 17, 189-255 (1985)]. Fifty percent of patients with therapy related AML show chromosome 5 abnormalities (interstitial deletions or monosomy) and cytogenetic analysis of the deletions has revealed that one segment, consisting of bands

5q23-31, is absent in the malignant cells of all patients who have aberrations of chromosome 5. These data suggest that loss of a critical DNA sequence leading to hemizygosity (or homozygosity) of a recessive allele may play an important role in the pathogenesis of these disorders, a mechanism substantiated for retinoblastoma. Although genes for a number of growth factors and receptors (IL-3, GM-CSF, β_2-adrenergic receptor, endothelial cell growth factor, CSF-1, c-fms, pDGF receptor) are clustered in or near this region, Egr-1 (by virtue of its zinc fingers) is the only member of this group with potential transcriptional regulatory activity. It is therefore possible that its absence could lead to deregulated cell growth.

EXAMPLE 3

In Vitro Expression of Mouse Egr-1 cDNA

A 2.1 kb ApaI/ApaI fragment (comprising nucleotides 120-2224 of Figure 1A) was isolated from the OC3.1 DNA clone. This fragment includes the translation start (ATG) codon at nucleotide position 259 designated in Figure 1A. The fragment was blunt-ended with T4 DNA polymerase and cloned into the Bluescript vector KS M13(+) containing a T3/T7 bacteriophage promoter. The (T3) sense transcript was generated and in vitro translated in a standard rabbit reticulocyte lysate system (Promega Biotec, Madison, WI. 53711) including ^{35}S methionine as a radiolabel. An analogous in vitro transcription system was developed using a BglII/BglII fragment of OC3.1 (including nucleotides 301-1958 and not including the translation start designated in Figure 1A). The T7 sense transcript was employed in the translation system. Differential characterization of translation products by autoradiographic SDS PAGE indicated that the ATG at nucleotide position 259 is preferred as

a translation start codon when all potential start sites are present.

EXAMPLE 4

Preparation of Antibodies:

A first synthetic peptide based on the sequence of amino acid residues 416-427 of mouse Egr-1 was prepared and provided with a carboxy terminal cysteine residue. The peptide, H-L-R-Q-K-D-K-K-A-D-K-S-C, was coupled to KLH and employed to immunize New Zealand white rabbits. Animals were initially immunized with 100 µg of the immunogen in Freund's Complete Adjuvant and every two weeks were boosted with 100 µg of immunogen with Freund's Incomplete Adjuvant. Sera, designated VPS10, were isolated after 68 days and displayed an antibody titer of 1:12,800 based on reactivity with the antigen used to prepare the anti-sera.

A second synthetic peptide, based on residues 399 to 415 of mouse Egr-1, was prepared. The peptide, C-G-R-K-R-A-R-S-D-E-R-K-R-H-T-K-I, was coupled to KLH and used to immunize rabbits as above, resulting in the production of antisera (designated VPS2) with a titer of 1:400.

EXAMPLE 5

Isolation of Genomic Mouse Egr-1 Clone and Characterization of Regulatory Regions

A mouse Balb/c 3T3 genomic library was prepared in a Stratagene (La Jolla, California) vector, λFIX, according to the manufacturer's instructions and probed using 1% SDS, 1 M NaCl, and 10% dextran sulfate at 65°C with stringent final wash in 0.2 x SSC at 65°C with a 2.1 kb ApaI/ApaI fragment and a 3.1 kb Eco RI/Eco

RI fragment derived from digestion of pUC13 including the mouse Egr-1 clone OC3.1. One positive clone, from approximately 300,000 screened, was designated mgEgr-1.1 and also hybridized to the extreme 5'-end 120 bp Eco RI-Apa I fragment from plasmid OC3.1.

A 2.4 kb Pvu-II-PvuII fragment and a 6.6 kb XbaI-XbaI fragment (see Figure 5) derived from the mgEgr-1.1 clone were subcloned into the SmaI and XbaI sites of pUC13 and pUC18 respectively, and the resulting plasmids (designated as p2.4 and p6.6) were used for restriction mapping analysis of transcription initiation sites and for nucleotide sequencing. Marked in Figure 4, and listed in Table 1, are possible regulatory elements identified in the 5' flanking sequence of mgEgr-1.1. A putative TATA motif (AAATA) is located 26 nucleotides upstream of the transcription start site. A "CCAAT" type sequence starts at nucleotide -337. Five different regions, each 10 nucleotides in length, located at -110, -342, -358, -374, and -412, are nearly identical to the inner core of the c-fos serum response element (Treisman, R., Cell, 46, 567 (1986)). Each has a 5-6 nucleotide AT rich stretch and is surrounded by the dinucleotide CC on the 5' side and GG on the other. Two potential TPA responsive elements (Lee, W., et al., Cell, 49, 741 (1987) and Angel, P., et al., Cell, 49, 729 (1987)) are located at nucleotides -610 and -867. Four consensus Spl (Briggs, M.R., et al., Science, 234, 47 (1986) binding sequences are at position -285, -649, -700 and -719. In addition, two sequences have been identified that might serve as cAMP response elements (Montimy, M.R., et al., Nature, 328, 175 (1987)) (-138 and -631).

TABLE 1

Location and Identification of Potential Regulatory Elements

Element	Sequence[1]	Location[2]
TATA	AAATA	-26 to -22
CCAAT	CCAAT	-337 to -333
Serum Response Element Consensus		
GATGTCCATATTAGGACATC	TCCTTCCATATTAGGGCTTC	-110 to -91
CC TA AT GG	GTGGCCC-AATATGGCCCTG	-342 to -324
G C	CAGCGCCTTATATGGAGTGG	-358 to -339
	ACAGACCTTATTTGGGCAGC	-374 to -355
	AAACGCCATATAAGGAGCAG	-412 to -393
TPA Responsive Element (AP1 binding site) Consensus		
C C	CTGACTCG	-610 to -603
TGACT A	CTGACTGG	-867 to -860
G A		
Sp1 binding site	GGGCGG	-285 to -280
	GGGCGG	-649 to -644
	CCGCCC	-700 to -695
	GGGCGG	-719 to -714
cAMP Response Element Consensus		
TGACGTCA	TCACGTCA	-138 to -131
	TGACGGCT	-631 to -624

1. The underlined bases in the mouse Egr-1 gene sequence are those that do not match the consensus sequence.

2. The location numbers refer to the nucleotides of the mouse Egr-1 gene as indicated in Figure 4.

To obtain the genomic sequence and the intron-exon gene structure, specific oligonucleotides (17-mers at positions 83, 122, 174, 200, 379, 543, 611, 659, 905, 920, 1000, 1200, 1400, 1600, 1800, 2100, 2353, 2650, 2825) of the OC3.1 cDNA sequence (see Figure 1A) were used as primers for double stranded sequencing of plasmids p2.4 and p6.6. Comparison of the Egr-1 genomic sequence to the Egr-1 cDNA sequence showed the Egr-1 gene consists of 2 exons and a single 700 bp intron (between nucleotide position 556 and 557 as numbered in Figure 1A and as shown in Figure 5). Both the 5' and 3' splice junction sequences (not shown) are in excellent agreement with the consensus boundary sequences. Mount, S.M., Nucleic Acids Res., 10, 459 (1982).

EXAMPLE 6

Isolation and Characterization of Human EGR2 cDNA

A human genomic placental library in the vector EMBL3, prepared by Dr. C. Westbrook of the University of Chicago according to procedures described in Frischauff et al., Jour. Mol. Biol., 170, 827-842 (1983), and a human leukocyte cosmid library prepared according to procedures described in Proc. Nat'l. Acad. Sci. (USA), 80, 5225-5229 (1983), were probed with the 2.1 kb ApaI fragment of OC3.1 (described in Example 5) using 1% SDS, 1 M NaCl and 10% dextrose sulfate at 50-55°C with a non-stringent final wash in 2 x SSC at 50-55°C. A single positive clone (designated HG6) was isolated from the first library and four clones (designated HG17, 18, 19 and 21, respectively) were isolated from the second library. A 6.6 kb SalI/EcoRI fragment of clone HG6 was found to hybridize with a 332 base pair HpaII/HpaII fragment of the mouse Egr-1 gene,

which letter fragment spans the putative zinc finger region. The 6.6 kb fragment, in turn, was employed to probe a cDNA library derived from human fibroblasts which have been stimulated for three hours with 20% fetal calf serum in the presence of 10 µg/ml cyclohexamide. About 10,000 clones were screened and the fifty positive clones obtained (designated "zap-1 through zap-50") are being subjected to nucleotide sequence analysis. Preliminary sequence analysis reveals that three clones, zap-2, zap-8, and zap-32, all encode the same transcript, namely a protein designated human EGR2, shown in Figure 3. Preliminary analysis indicates approximately 92% homology between mouse Egr-1 and human EGR2 polypeptides in the zinc finger regions, but substantially less homology in the amino and carboxy terminal regions. Chromosome mapping studies, similar to those described in Example 2, indicate that human chromosome 10, at bands q21-22, constitutes a locus for the human EGR2 gene.

The plasmid zap-32, containing the full length human EGR2 clone, was used as a probe in Southern blot analysis on DNAs from 58 unrelated Caucasians. It was found that Hind III detects a simple two-allele polymorphism with bands at either 8.0 kb (A1) or 5.6 kb and 2.4 kb (A2). No constant bands were detected. The frequency of A1 was 0.90 and that of A2 was 0.10. No polymorphisms were detected for Apa I, BamH I, Ban II, Bgl I, Bgl II, BstE II, Dra I, EcoR I, EcoR V, Hinc II, Msp I, Pst I, Pvu II, Rsa I, Sac I, and Taq I in 10 unrelated individuals. Co-dominant segregation of the Hind III RFLP was observed in four large kindreds with a total of more than 350 individuals.

These data will be useful in gene linkage studies for mapping genes for certain genetic disorders. For example, the gene responsible for the dominantly inherited syndrome, multiple endocrine

neoplasia, type 2A (MEN-2A) has been assigned by linkage to chromosome 10. Simpson, et al., Nature, 328, 528 (1987). Studies are currently underway to determine the linkage relationship between MEN-2A and EGR2 and are expected to be useful in cloning the MEN-2A gene as well as in serving as a diagnostic marker for the disease.

EXAMPLE 7

Recombinant Expression Of Fusion Proteins

A 322 base HpaII/HpaII fragment (comprising nucleotides 1231-1553) derived from the OC3.1 cloned DNA was treated with DNA polymerase to fill in the single stranded ends. This fragment was inserted in plasmid pEX3 (obtained from K. Stanley, European Molecular Biology Laboratory, Postfach 10.2209, 6900 Heidelberg, F.R.G.) digested with SmaI. Stanley, K.K., et al., EMBO J., 3, 1429 (1984). This insertion placed the Egr-1 encoding DNA fragment in the same reading frame as plasmid DNA encoding cro-β-galactosidase, allowing for the expression of a fusion protein comprising the amino terminal residues of cro-β-galactosidase and 108 residues of Egr-1 amino acids 325 to 432. This cro-β-galactosidase/Egr-1 fusion plasmid, designated pFIG, was used to transform E. coli NF1.

Induced (42°C) and un-induced (30°C) cultured cell lysates from growth of the transformed NF1 cells were then analyzed by SDS-PAGE. Upon Coomassie stain analysis, only induced cell lysates included an approximately 108 kd product, indicating presence of the projected expression product. Western blot analysis, using the rabbit polyclonal anti-peptide antibody VPS10 (see Example 4) raised against H-L-R-Q-K-D-K-K-A-D-K-S-C, confirmed that the fusion protein product contained Egr sequences.

In a separate construction, a mouse Egr-1 insert, from plasmid 0C3.1, was fused, in frame, to a plasmid containing sequences from bovine growth hormone according to the methods described in Slamon, D.J., et al., Science, 233, 347 (1986). The resultant plasmid, designated pV4, comprised a fusion protein containing a fusion gene coding for bovine growth hormone amino acids 1 to 192 and Egr-1 amino acids 2 to 533. This bGH/mouse Egr-1 DNA fusion plasmid, designated pV4, was expressed in E. coli and the resulting fusion protein, designated V4, was identified in Western blots by its reactivity with a bGH monoclonal antibody and its reactivity with VPS10 rabbit anti-Egr-1 peptide antiserum, prepared according to Example 4.

EXAMPLE 8

Determination of Egr Levels in Human Tumor and Non-Tumor Tissue

Using the mouse Egr-1 0C68 probe, Northern blot analyses were conducted to determine the levels of transcription of Egr protein encoding DNA in tumor versus surrounding normal tissue from resected human tumor specimens. The tumor samples were from lung (12), colon (7), colon mesastasis (1), bladder (1), rectal (1), giant cell (1), hepatoma (1), breast (1), MFH (malignant fibrous histiocytoma) (1), osteosarcoma (1) and rhabdomyosarcoma (1). In about 50% of these cases, there is markedly decreased (about three to ten-fold) expression of the Egr mRNA in tumor versus normal tissue. One implication of this finding is that Egr proteins of the invention may function as part of a negative regulatory pathway. In any event, it is clear that DNA sequences and antibodies of the invention are susceptible to use in differential diagnoses between tumorous and non-tumorous cell types.

It will be apparent from consideration of the foregoing illustrative examples that the present invention constitutes a substantial advance in the art and the achievement of a major goal in molecular biology, i.e., the characterization of genes which play a regulatory role in mammalian cell proliferation and differentiation. It will thus be understood that the information provided herein constitutes a basis for straightforward development of useful methods and materials not specifically the subject of the above examples. By way of illustration, possession of knowledge concerning the base sequence of cDNA and genomic DNA sequences encoding distinct mouse Egr-1 and human EGR2 early growth regulatory proteins comprising histidine-cysteine zinc finger amino acid sequences makes possible the isolation of other such structurally related proteins. The substantial homology between the zinc finger regions of Egr-1 and EGR2 coupled with lack of homology in other protein regions, when considered in light of the ability of Egr-1 probes to localize to human chromosome 5 while EGR2 probes localize to human chromosome 10, essentially assures the straightforward isolation of a human gene (provisionally designated "human EGR1") which encodes a protein more closely homologous to Egr-1 and a mouse gene (Egr-2) encoding a protein more closely homologous to EGR2.

While the above examples provide only limited illustration of <u>in vitro</u> and <u>in vivo</u> expression of DNA sequences of the invention, known recombinant techniques are readily applicable to development of a variety of procaryotic and eucaryotic expression systems for the large scale production of Egr proteins and even development of gene therapy regimens.

Knowledge of the specifically illustrated mouse Egr-1 and human EGR2 proteins of the invention has been demonstrated to provide a basis for preparation of

highly useful antibodies, also provides a wealth of information concerning the nature of protein-nucleic acid interactions which, in turn, constitutes a basis for determination of significant early growth regulatory events. For example, and by analogy to steroid receptor protein structures, analysis of the structure of regions flanking the zinc fingers of Egr-1 and EGR2 and related proteins of the invention is expected to allow for identification of substances which may interact with the proteins to alter their DNA interactive capacities and thus provide the basis for inhibition or augmentation of their regulatory functions. Moreover, information available concerning specific events of DNA interaction of Egr proteins of the invention will permit, e.g., identification and use of potential competitive inhibitors of these proteins.

Just as Egr encoding DNA of the invention is conspicuously susceptible to use in differentiation of human tumor and non-tumor cells, antibodies prepared according to the invention are expected to be useful in differential screening of cells based on relative cellular concentrations of mRNA expression products and in the determination of specific genes susceptible to regulation by such products.

Because numerous modifications and variations in the practice of the present invention are expected to occur to those skilled in the art, only such limitations as appear in the appended claims should be placed thereon.

WHAT IS CLAIMED IS

1. A purified and isolated DNA sequence encoding a mammalian early growth regulatory protein which comprises one or more histidine-cysteine zinc finger amino acid sequences.

2. The DNA sequence according to claim 1 encoding human EGR2 protein.

3. The DNA sequence according to claim 1 encoding mouse Egr-1 protein.

4. The DNA sequence according to claim 1 which is a cDNA sequence.

5. The DNA sequence according to claim 1 which is a genomic DNA sequence.

6. The DNA sequence according to claim 1 which is a wholly or partially synthetic DNA sequence.

7. The DNA sequence according to claim 1 operatively associated with an homologous or heterologous expression control DNA sequence.

8. The DNA sequence according to claim 1 selected from the group consisting of the DNA sequences set out in Figures 1A, 3, and 4.

9. A procaryotic or eucaryotic host cell transformed or transfected with a DNA sequence comprising a DNA sequence according to claim 1 or 7.

10. A viral or circular DNA plasmid vector comprising a DNA sequence according to claim 1.

11. The viral or circular DNA plasmid vector according to claim 10 further comprising an expression control DNA sequence operatively associated with said early growth regulatory protein encoding DNA.

12. A method for the production of an early growth regulatory protein comprising:

growing, in culture, a host cell transformed or transfected with a DNA sequence according to claim 1; and

isolating from said host cell or culture the polypeptide product of the expression of said DNA sequence.

13. A method for the production of an early growth regulatory protein comprising:

disposing a DNA sequence according to claim 1 in a cell free transcription and translation system; and

isolating from said system the polypeptide product of the expression of said DNA sequence.

14. The polypeptide product of the in vitro or in vivo expression of part or all of a protein encoding region of a DNA sequence according to claim 1.

15. The polypeptide product according to claim 14 which is a fusion protein comprising part or all of a mammalian early growth regulatory protein which comprises one or more histidine-cysteine zinc finger amino acid sequences and part or all of a heterologous proteins.

16. The polypeptide product according to claim 15, which comprises a fusion of cro-β-galactosidase and Egr-1 amino acid sequences or bovine growth hormone and Egr-1 amino acid sequences.

17. A purified and isolated mammalian early growth regulatory protein comprising one or more histidine-cysteine zinc finger amino acid sequences.

18. The protein according to claim 17 and having the amino acid sequence set out in Figure 1A.

19. The protein according to claim 17 and having the amino acid sequence set out in Figure 3.

20. A synthetic peptide duplicative of a sequence of amino acids present in a mammalian early growth regulatory protein comprising one or more histidine-cysteine zinc finger amino acid sequences and sharing at least one antigenic epitope of such protein.

21. The synthetic peptide of claim 20 and having an amino acid sequence partially duplicative of the amino acid sequence set out in Figure 1A.

22. The synthetic peptide of claim 20 and having an amino acid sequence partially duplicative of the amino acid sequence set out in Figure 3.

23. The synthetic peptide of claim 20 and further characterized as duplicative of an amino acid sequence not involved in DNA binding functions.

24. An antibody specifically immunoreactive with at least one epitope of a mammalian early growth regulatory protein comprising one or more histidine-cysteine zinc finger amino acid sequences.

25. The antibody according to claim 24 wherein said epitope is other than an epitope within the DNA binding functional region thereof.

26. The antibody according to claim 24 selected from the group consisting of monoclonal and polyclonal antibodies.

27. The antibody according to claim 24 capable of specifically binding with a proteinaceous material comprising an amino acid sequence duplicating an antigenic epitope within the following amino acid sequence:

H-L-R-Q-K-D-K-K-A-D-K-S-C.

28. The antibody according to claim 24 capable of specifically binding with a proteinaceous material comprising an amino acid sequence duplicating an antigenic epitope within the following amino acid sequence:

C-G-R-K-F-A-R-S-D-E-R-K-R-H-T-K-I.

29. A method for quantitative detection of a mammalian early growth regulatory protein comprising one or more histidine-cysteine zinc finger amino acid sequences based on the immunological reaction of the same with an antibody according to claim 24.

30. A method for quantitative detection within a sample of messenger RNA transcripts for mammalian early growth regulatory proteins comprising one or more histidine-cysteine zinc finger amino acid sequences comprising the step of hybridizing RNA within said sample with a DNA sequence partially or wholly duplicating a DNA sequence according to claim 1.

31. A method for quantitative detection within a sample of DNA encoding for mammalian early growth regulatory proteins comprising one or more histidine-cysteine zinc finger amino acid sequences comprising the step of hybridizing DNA within said sample with a DNA sequence partially or wholly duplicating a DNA sequence according to claim 1.

32. A method for detecting a disease genetically linked to a mammalian Egr gene comprising the step of quantitating mammalian early growth regulatory DNA sequences according to claim 1.

```
        10          20          30          40          50          60
GGGGAGCCGCCGCCGCGATTCGCCGCCGCCGCCAGCTTCCGCCGCCGCAAGATCGGCCCC

        70          80          90         100         110         120
TGCCCCAGCCTCCGCGGCAGCCCTGCGTCCACCACGGGCCGCGGCTACCGCCAGCCTGGG

       130         140         150         160         170         180
GGCCCACCTACACTCCCCGCAGTGTGCCCCTGCACCCCGCATGTAACCCGGCCAACCCCC

       190         200         210         220         230         240
GGCGAGTGTGCCCTCAGTAGCTTCGGCCCCGGGCTGCGCCCACCACCCAACATCAGTTCT

       250         260         270         280         290         300
CCAGCTCGCTGGTCCGGGATGGCAGCGGCCAAGGCCGAGATGCAATTGATGTCTCCGCTG
                  MetAlaAlaAlaLysAlaGluMetGlnLeuMetSerProLeu

       310         320         330         340         350         360
CAGATCTCTGACCCGTTCGGCTCCTTTCCTCACTCACCCACCATGGACAACTACCCCAAA
GlnIleSerAspProPheGlySerPheProHisSerProThrMetAspAsnTyrProLys

       370         380         390         400         410         420
CTGGAGGAGATGATGCTGCTGAGCAACGGGGCTCCCCAGTTCCTCGGTGCTGCCGGAACC
LeuGluGluMetMetLeuLeuSerAsnGlyAlaProGlnPheLeuGlyAlaAlaGlyThr
```

FIGURE 1.1

```
       430        440        450        460        470        480
CCAGAGGGCAGCGGCGGTAATAGCAGCAGCAGCACCAGCAGCGGGGGCGGTGGTGGGGGC
ProGluGlySerGlyGlyAsnSerSerSerSerThrSerSerGlyGlyGlyGlyGlyGly

       490        500        510        520        530        540
GGCAGCAACAGCGGCAGCAGCGCCTTCAATCCTCAAGGGGAGCCGAGCGAACAACCCTAT
GlySerAsnSerGlySerSerAlaPheAsnProGlnGlyGluProSerGluGlnProTyr

       550        560        570        580        590        600
GAGCACCTGACCACAGAGTCCTTTTCTGACATCGCTCTGAATAATGAGAAGGCGATGGTG
GluHisLeuThrThrGluSerPheSerAspIleAlaLeuAsnAsnGluLysAlaMetVal

       610        620        630        640        650        660
GAGACGAGTTATCCCAGCCAAACGACTCGGTTGCCTCCCATCACCTATACTGGCCGCTTC
GluThrSerTyrProSerGlnThrThrArgLeuProProIleThrTyrThrGlyArgPhe

       670        680        690        700        710        720
TCCCTGGAGCCCCGCACCCAACAGTGGCAACACTTTGTGGCCTGAACCCCTTTTCAGCCTA
SerLeuGluProAlaProAsnSerGlyAsnThrLeuTrpProGluProLeuPheSerLeu

       730        740        750        760        770        780
GTCAGTGGCCTCGTGAGCATGACCAATCCTCCGACCTCTTCATCCTCGGCGCCTTCTCCA
ValSerGlyLeuValSerMetThrAsnProProThrSerSerSerSerAlaProSerPro

       790        800        810        820        830        840
GCTGCTTCATCGTCTTCCTCTGCCTCCCAGAGCCCGCCCCTGAGCTGTGCCGTGCCGTCC
AlaAlaSerSerSerSerSerAlaSerGlnSerProProLeuSerCysAlaValProSer
```

FIGURE 1.2

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      850        860        870        880        890        900
AACGACAGCAGTCCCATCTACTCGGCTGCGCCCACCTTTCCTACTCCCAACACTGACATT
AsnAspSerSerProIleTyrSerAlaAlaProThrPheProThrProAsnThrAspIle
      910        920        930        940        950        960
TTTCCTGAGCCCCAAAGCCAGGCCTTTCCTGGCTCGGCAGGCACAGCCTTGCAGTACCCG
PheProGluProGlnSerGlnAlaPheProGlySerAlaGlyThrAlaLeuGlnTyrPro

      970        980        990       1000       1010       1020
CCTCCTGCCTACCCTGCCACCAAAGGTGGTTTCCAGGTTCCCATGATCCCTGACTATCTG
ProProAlaTyrProAlaThrLysGlyGlyPheGlnValProMetIleProAspTyrLeu

     1030       1040       1050       1060       1070       1080
TTTCCACAACAACAGGGAGACCTGAGCCTGGGCACCCCAGACCAGAAGCCCTTCCAGGGT
PheProGlnGlnGlnGlyAspLeuSerLeuGlyThrProAspGlnLysProPheGlnGly

     1090       1100       1110       1120       1130       1140
CTGGAGAACCGTACCCAGCAGCCTTCGCTCACTCCACTATCCACTATTAAAGCCTTCGCC
LeuGluAsnArgThrGlnGlnProSerLeuThrProLeuSerThrIleLysAlaPheAla

     1150       1160       1170       1180       1190       1200
ACTCAGTCGGGCTCCCAGGACTTAAAGGCTCTTAATACCACCTACCAATCCCAGCTCATC
ThrGlnSerGlySerGlnAspLeuLysAlaLeuAsnThrThrTyrGlnSerGlnLeuIle

     1210       1220       1230       1240       1250       1260
AAACCCAGCCGCATGCGCAAGTACCCCAACCGGCCCAGCAAGACACCCCCCCATGAACGC
LysProSerArgMetArgLysTyrProAsnArgProSerLysThrProProHisGluArg
```

FIGURE 1.3

```
      1270       1280       1290       1300       1310       1320
CCATATGCTTGCCCTGTCGAGTCCTGCGATCGCCGCTTTTCTCGCTCGGATGAGCTTACC
ProTyrAlaCysProValGluSerCysAspArgArgPheSerArgSerAspGluLeuThr

      1330       1340       1350       1360       1370       1380
CGCCATATCCGCATCCACACAGGCCAGAAGCCCTTCCAGTGTCGAATCTGCATGCGTAAC
ArgHisIleArgIleHisThrGlyGlnLysProPheGlnCysArgIleCysMetArgAsn

      1390       1400       1410       1420       1430       1440
TTCAGTCGTAGTGACCACCTTACCACCCACATCCGCACCCACACAGGCGAGAAGCCTTTT
PheSerArgSerAspHisLeuThrThrHisIleArgThrHisThrGlyGluLysProPhe

      1450       1460       1470       1480       1490       1500
GCCTGTGACATTTGTGGGAGGAAGTTTGCCAGGAGTGATGAACGCAAGAGGCATACCAAA
AlaCysAspIleCysGlyArgLysPheAlaArgSerAspGluArgLysArgHisThrLys

      1510       1520       1530       1540       1550       1560
ATCCATTTAAGACAGAAGGACAAGAAAGCAGACAAAAGTGTGGTGGCCTCCCCGGCTGCC
IleHisLeuArgGlnLysAspLysLysAlaAspLysSerValValAlaSerProAlaAla

      1570       1580       1590       1600       1610       1620
TCTTCACTCTCTTCTTACCCATCCCCAGTGGCTACCTCCTACCCATCCCCTGCCACCACC
SerSerLeuSerSerTyrProSerProValAlaThrSerTyrProSerProAlaThrThr

      1630       1640       1650       1660       1670       1680
TCATTCCCATCCCCTGTGCCCACTTCCTACTCCTCTCCTGGCTCCTCCACCTACCCATCT
SerPheProSerProValProThrSerTyrSerSerProGlySerSerThrTyrProSer
```

FIGURE 1.4

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      1690        1700        1710        1720        1730        1740
CCTGCGCACAGTGGCTTCCCGTCGCCGTCAGTGGCCACCACCTTTGCCTCCGTTCCACCT
ProAlaHisSerGlyPheProSerProSerValAlaThrThrPheAlaSerValProPro

      1750        1760        1770        1780        1790        1800
GCTTTCCCCACCCAGGTCAGCAGCTTCCCGTCTGCGGGCGTCAGCAGCTCCTTCAGCACC
AlaPheProThrGlnValSerSerPheProSerAlaGlyValSerSerSerPheSerThr

      1810        1820        1830        1840        1850        1860
TCAACTGGTCTTTCAGACATGACAGCGACCTTTTCTCCCAGGACAATTGAAATTTGCTAA
SerThrGlyLeuSerAspMetThrAlaThrPheSerProArgThrIleGluIleCys

      1870        1880        1890        1900        1910        1920
AGGGAATAAAAGAAAGCAAAGGGAGAGGCAGGAAAGACATAAAAGCACAGGAGGGAAGAG

      1930        1940        1950        1960        1970        1980
ATGGCCGCAAGAGGGGCCACCTCTTAGGTCAGATGGAAGATCTCAGAGCCAAGTCCTTCT

      1990        2000        2010        2020        2030        2040
ACTCACGAGTAGAAGGACCGTTGGCCAACAGCCCTTTCACTTACCATCCCTGCCTCCCCC

      2050        2060        2070        2080        2090        2100
GTCCTGTTCCCTTTGACTTCAGCTGCCTGAAACAGCCATGTCCAAGTTCTTCACCTCTAT
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FIGURE 1.5

```
      2110        2120        2130        2140        2150        2160
CCAAAGGACTTGATTTGCATGGTATTGGATAAATCATTTCAGTATCCTCTCCATCACATG

      2170        2180        2190        2200        2210        2220
CCTGGCCCTTGCTCCCTTCAGCGCTAGACCATCAAGTTGGCATAAAGAAAAAAAAATGGG

      2230        2240        2250        2260        2270        2280
TTTGGGCCCTCAGAACCCTGCCCTGCATCTTTGTACAGCATCTGTGCCATGGATTTTGTT
      2290        2300        2310        2320        2330        2340
TTCCTTGGGGTATTCTTGATGTGAAGATAATTTGCATACTCTATTGTATTATTTGGAGTT

      2350        2360        2370        2380        2390        2400
AAATCCTCACTTTGGGGGAGGGGGGAGCAAAGCCAAGCAAACCAATGATGATCCTCTATT

      2410        2420        2430        2440        2450        2460
TTGTGATGACTCTGCTGTGACATTAGGTTTGAAGCATTTTTTTTTTTCAAGCAGCAGTCCT

      2470        2480        2490        2500        2510        2520
AGGTATTAACTGGAGCATGTGTCAGAGTGTTGTTCCGTTAATTTTGTAAATACTGGCTCG
```

FIGURE 1.6

```
2530      2540      2550      2560      2570      2580
ACTGTAACTCTCACATGTGACAAAGTATGGTTTGTTTGGGTTTTTGTTTTGAGAAT

2590      2600      2610      2620      2630      2640
TTTTTTTGCCCGTCCCTTTGGTTTCAAAAGTTTCACGTCTTGGTGCCTTTTTGTGACACG

2650      2660      2670      2680      2690      2700
CCTTCCGATGGCTTGACATGCGCAGATGTGAGGGACACGCTCACCTTAGCCTTAAGGGGG

2710      2720      2730      2740      2750      2760
TAGGAGTGATGTGTTGGGGGAGGCTTGAGAGCAAAAACGAGGAAGAGGGCTGAGCTGAGC

2770      2780      2790      2800      2810      2820
TTTCGGTCTCCAGAATGTAAGAAGAAAAAATTTAAACAAAAATCTGAACTCTCAAAAGTC
```

FIGURE 1.7

```
2830      2840      2850      2860      2870      2880
TATTTTTCTAAACTGAAAATGTAAATTTATACATCTATTCAGGAGTTGGAGTGTTGTGGT

2890      2900      2910      2920      2930      2940
TACCTACTGAGTAGGCTGCAGTTTTTGTATGTTATGAACATGAAGTTCATTATTTTGTGG

2950      2960      2970      2980      2990      3000
TTTTATTTTTACTTTGTACTTGTTTGCTTAAACAAGTAACCTGTTTGGCTTATAAACA

3010      3020      3030      3040      3050      3060
CATTGAATGCGCTCTATTGCCCATGGGATATGTGGTGTATCCTTCAGAAAAATTAAAA

3070      3080
GGAAAAATAAAAAAAAAAAAAA
```

FIGURE 1.8

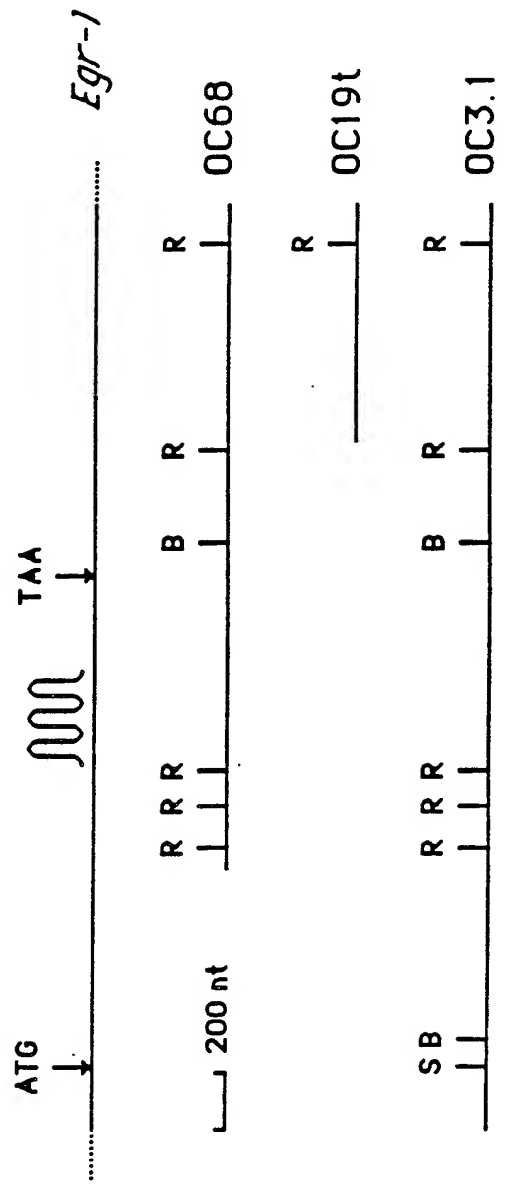

FIGURE 1B

"zinc finger" consensus sequence	T	G	E	R/K	P	F/Y	X	C	X	X	X	X	C	X	X	X	F	X	X	X	X	L	X	X	H	X	X	X	H		
Murine Egr-1	P	H	E	R	P	Y	A	C	P	V	E	S	C	D	R	R	F	S	R	S	D	E	L	T	R	H	I	R	I	H	
	T	G	Q	K	P	F	Q	C	-	-	R	I	C	M	R	N	F	S	R	S	D	H	L	T	T	H	I	R	T	H	
	T	G	E	K	P	F	A	C	-	-	D	I	C	G	R	K	F	A	R	S	D	E	R	K	R	H	T	K	I	H	
Drosophila Kruppel	S	R	D	K	S	F	T	C	K	I	-	-	C	S	R	S	F	G	Y	K	H	V	L	Q	N	H	E	R	T	H	
	T	G	E	K	P	F	E	C	P	E	-	-	C	D	K	R	F	T	R	D	H	H	L	K	T	H	M	R	L	H	
	T	G	E	K	P	Y	H	C	S	H	-	-	C	D	R	Q	F	V	Q	V	A	N	L	R	R	H	L	R	V	H	
	T	G	E	R	P	Y	T	C	E	I	-	-	C	D	G	K	F	S	D	S	N	Q	L	K	S	H	M	L	V	H	
Xenopus TFIIIA (finger 2)	T	G	E	K	P	F	P	C	K	E	E	G	C	E	K	G	F	T	S	L	H	H	L	T	R	H	S	L	T	H	

FIGURE 2

```
        10         20         30         40         50         60
TTTTTTTTTTTGGTGTGTGTGGTGGTTGTTTTTAAGTGTGGAGGGCAAAAGGAGATACCA

        70         80         90        100        110        120
TCCCAGGCTCAGTCCAACCCCTCTCCAAAACNGTGTCTTTTCTGACACTCCAGGTAGCGA

       130        140        150        160        170        180
GGGAGTTGGGTCTCCAGGTTGTGCGAGGAGCAAATGATGACCGCCAAGGCCGTAGACAAA
                                      MetMetThrAlaLysAlaValAspLys

       190        200        210        220        230        240
ATCCCAGTAACTCTCAGTGGTTTTGTGCACCAGCTGTCTGACAACATCTACCCGGTGGAG
IleProValThrLeuSerGlyPheValHisGlnLeuSerAspAsnIleTyrProValGlu

       250        260        270        280        290        300
GACCTCGCCGCCACGTCGGTGACCATCTTTCCCAATGCCGAACTGGGAGGCCCCTTTGAC
AspLeuAlaAlaThrSerValThrIlePheProAsnAlaGluLeuGlyGlyProPheAsp

       310        320        330        340        350        360
CAGATGAACGGAGTGGCCGGAGATGGCATGATCAACATTGACATGACTGGAGAGAAGAGG
GlnMetAsnGlyValAlaGlyAspGlyMetIleAsnIleAspMetThrGlyGluLysArg
```

FIGURE 3.1

```
       370        380        390        400        410        420
TCGTTGGATCTCCCATATCCCAGCAGCTTTGCTCCCGTCTCTGCACCTAGAAACCAGACC
SerLeuAspLeuProTyrProSerSerPheAlaProValSerAlaProArgAsnGlnThr

       430        440        450        460        470        480
TTCACTTACATGGGCAAGTTCTCCATTGACCCACAGTACCCTGGTGCCAGCTGCTACCCA
PheThrTyrMetGlyLysPheSerIleAspProGlnTyrProGlyAlaSerCysTyrPro

       490        500        510        520        530        540
GAAGGCATAATCAATATTGTGAGTGCAGGCATCTTGCAAGGGGTCACTTCCCCAGCTTCA
GluGlyIleIleAsnIleValSerAlaGlyIleLeuGlnGlyValThrSerProAlaSer

       550        560        570        580        590        600
ACCACAGCCTCATCCAGCGTCACCTCTGCCTCCCCCAACCCACTGGCCACAGGACCCCTG
ThrThrAlaSerSerSerValThrSerAlaSerProAsnProLeuAlaThrGlyProLeu

       610        620        630        640        650        660
GGTGTGTGCACCATGTCCCAGACCCAGCCTGACCTGGACCACCTGTACTCTCCGCCACCG
GlyValCysThrMetSerGlnThrGlnProAspLeuAspHisLeuTyrSerProProPro

       670        680        690        700        710        720
CCTCCTCCTCCTTATTCTGGCTGTGCAGGAGACCTCTACCAGGACCCTTCTGCGTTCCTG
ProProProProTyrSerGlyCysAlaGlyAspLeuTyrGlnAspProSerAlaPheLeu
```

FIGURE 3.2

```
      730        740        750        760        770        780
TCAGCAGCCACCACCTCCACCTCTTCCTCTCTGGCCTACCCACCACCTCCTTCCTATCCA
SerAlaAlaThrThrSerThrSerSerSerLeuAlaTyrProProProProSerTyrPro

      790        800        810        820        830        840
TCCCCCAAGCCAGCCACGGACCCAGGTCTCTTCCCAATGATCCCAGACTATCCTGGATTC
SerProLysProAlaThrAspProGlyLeuPheProMetIleProAspTyrProGlyPhe

      850        860        870        880        890        900
TTTCCATCTCAGTGCCAGAGAGACCTACATGGTACAGCTGGCCCAGACCGTAAGCCCTTT
PheProSerGlnCysGlnArgAspLeuHisGlyThrAlaGlyProAspArgLysProPhe

      910        920        930        940        950        960
CCCTGCCCACTGGACACCCTGCGGGTGCCCCCTCCACTCACTCCACTCTCTACAATCCGT
ProCysProLeuAspThrLeuArgValProProProLeuThrProLeuSerThrIleArg

      970        980        990       1000       1010       1020
AACTTTACCCTGGGGGGCCCCAGTGCTGGGATGACCGGACCAGGGGCCAGTGGAGGCAGC
AsnPheThrLeuGlyGlyProSerAlaGlyMetThrGlyProGlyAlaSerGlyGlySer

     1030       1040       1050       1060       1070       1080
GAGGGACCCCGGCTGCCTGGTAGCAGCTCAGCAGCAGCAGCAGCCGCCGCCGCCGCCGCC
GluGlyProArgLeuProGlySerSerSerAlaAlaAlaAlaAlaAlaAlaAlaAlaAla
```

FIGURE 3.3

```
      1090       1100       1110       1120       1130       1140
TATAACCCACACCACCTGCCACTGCGGCCCATTCTGAGGCCTCGCAAGTACCCCAACAGA
TyrAsnProHisHisLeuProLeuArgProIleLeuArgProArgLysTyrProAsnArg

      1150       1160       1170       1180       1190       1200
CCCAGCAAGACGCCGGTGCACGAGAGGCCCTACCCGTGCCCAGCAGAAGGCTGCGACCGG
ProSerLysThrProValHisGluArgProTyrProCysProAlaGluGlyCysAspArg

      1210       1220       1230       1240       1250       1260
CGGTTCTCCCGCTCTGACGAGCTGACACGGCACATCCGAATCCACACTGGGCATAAGCCC
ArgPheSerArgSerAspGluLeuThrArgHisIleArgIleHisThrGlyHisLysPro

      1270       1280       1290       1300       1310       1320
TTCCAGTGTCGGATCTGCATGCGCAACTTCAGCCGCAGTGACCACCTCACCACCCATATC
PheGlnCysArgIleCysMetArgAsnPheSerArgSerAspHisLeuThrThrHisIle

      1330       1340       1350       1360       1370       1380
CGCACCCACACCGGTGAGAAGCCCTTCGCCTGTGACTACTGTGGCCGAAAGTTTGCCCGG
ArgThrHisThrGlyGluLysProPheAlaCysAspTyrCysGlyArgLysPheAlaArg

      1390       1400       1410       1420       1430       1440
AGTGATGAGAGGAAGCGCCACACCAAGATCCACCTGAGACAGAAAGAGCGGAAAAGCAGT
SerAspGluArgLysArgHisThrLysIleHisLeuArgGlnLysGluArgLysSerSer
```

FIGURE 3.4

```
      1450       1460       1470       1480       1490       1500
GCCCCCTCTGCATCGGTGCCAGCCCCCTCTACAGCCTCCTGCTCTGGGGGCGTGCAGGCC
AlaProSerAlaSerValProAlaProSerThrAlaSerCysSerGlyGlyValGlnAla

      1510       1520       1530       1540       1550       1560
TGGGGGTACCCTGTGCAGCAGTAACAGCAGCAGTCTTGGCGGAGGGCCGCTCGCCCCTTG
.TrpGlyTyrProValGlnGlnEnd

      1570       1580       1590       1600       1610       1620
CTCCTCTCGGACCCGGACACCTTGAGATGAGACTCAGGCTGATACACCAGCTCCCAAAGG

      1630       1640       1650       1660       1670       1680
TCCCGGAGGCCCTTTGTCCACTGGAGCTGCACAACAAACACTACCACCCTTTCCTGTCCC

      1690       1700       1710       1720       1730       1740
TCTCTCCCTTTGTTGGGCAAAGGGCTTTGGTGGAGCTAGCACTGCCCCCTTTCCACCTAG

      1750       1760       1770       1780       1790       1800
AAGCAGGTTCTTCCTAAAACTTAGCCCATTCTAGTCTCTCTTAGGTGAGTTGACTATCAA
```

FIGURE 3.5

```
      1810        1820        1830        1840        1850        1860
CCCAAGGCAAAGGGGAGGCTCAGAAGGAGGTGGTGTGGGGATCCCCTGGCCAAGAGGGCT

      1870        1880        1890        1900        1910        1920
GAGGTCTGACCCTGCTTTAAAGGGTTGTTTGACTAGGTTTTGCTACCCCACTTCCCCTTA

      1930        1940        1950        1960        1970        1980
TTTTGACCCATCACAGGTTTTTGACCCTGGATGTCAGAGTTGATCTAAGACGTTTTCTAC

      1990        2000        2010        2020        2030        2040
AATAGGTTGGGAGATGCTGATCCCTTCAAGTGGGGACAGCAAAAAGACAAGCAAAACTGA

      2050        2060        2070        2080        2090        2100
TGTGCACTTTATGGCTTGGGACTGATTTGGGGGACATTGTACAGTGAGTGAAGTATAGCC

      2110        2120        2130        2140        2150        2160
TTTATGCCACACTCTGTGGCCCTAAAATGGTGAATCAGAGCATATCTAGTTGTCTCAACC
```

FIGURE 3.6

```
2170      2180      2190      2200      2210      2220
CTTGAAGCAATATGTATTATACTCAGAGAACAGAAGTGCAATGTGATGGGAGGAACGT

2230      2240      2250      2260      2270      2280
AGCAATATCTGCTCCTTTTCGAGTTGTTTGAGAAATGTAGGCTATTTTTTTCAGTGTATAT

2290      2300      2310      2320      2330      2340
CCACTCAGATTTTTGTGTATTTTTTGATGTACCCACACTGTTCTCTAAATTCTGAATCTTTG

2350      2360      2370      2380      2390      2400
GGAAAAAATGTAAAGCATTTATGATCTCAGAGGTTAACTTATTTTAAGGGGGATGTACATA

2410      2420      2430      2440      2450      2460
TTCTCTGAAACTAGGATGCATGCAATTTGTGTTGGAAGTGTCCTTGGTCGCCTTGTGTGAT

2470      2480      2490      2500      2510      2520
GTAGACAAATGTTACAAGGCTGCATGTAAATGGGTTGCCTTATTATGGAGAAAAAAATCA
```

FIGURE 3.7

```
      2530        2540        2550        2560        2570        2580
CTCCCTGAGTTTAGTATGGCTGTATATTTATGCCTATTAATATTTCAAATTTTTTTTTAG

      2590        2600        2610        2620        2630        2640
AGTATATTTTTGTATGCTTTGTTTTGTGACTTAAAAGTGTTACCTTTGTAGTCAAATTTC

      2650        2660        2670        2680        2690        2700
AGATAAGAATGTACATAATGTTACCGGAGCTGANNNTTGTTTGGTCATTAGCTCTTAATA
      2710        2720        2730        2740        2750        2760
GTTGTGAAAAAATAAATCTATTCTAACGCAAAACCACTAACTGAAGTTCAGATATAATGG

      2770        2780        2790        2800        2810
ATGGTTTGTGACTATAGTGTAAATAAATACTTTTCAACAAAAAAAAAAAAAAAAAAA
```

FIGURE 3.8

-935
ACGGAGGGAA TAGCCTTTCG ATTCTGGGTG GTGCATTGGA AGCCCCCAGGC TCTANNACCC -876

-875
CCAACCTACT GACTGGTGGC CGAGTATGCA CCCGACTGCT AGCTAGGCAG TGTCCCAAGA -816

-815
ACCAGTAGCC AAATGTCTTG GCCTCAGTTT TCCCGGTGAC ACCTGGAAAG TGACCCTGCC -756

-755
ATTAGTAGAG GCTCAGGTCA GGGCCCCGCC TCTCCTGGGC GGCCTCTGCC CTAGCCCGCC -696

-695
CTGCCGCTCC TCCTCTCCGC AGGCTCGCTC CCACGGTCCC CGAGGTGGGC GGGTGAGCCC -636

-635
AGGATGACGG CTGTAGAACC CCGGCCTGAC TCGCCCTCGC CCCCGCGCCG GGCCTGGGCT -576

-575
TCCCTAGCCC AGCTCGCACC CGGGGGCCGT CGGAGCCGCC GCGCGCCCAG CTCTACGCGC -516

-515
CTGGCCCTCC CCACGCGGGC GTCCCCGACT CCCGCGCGCG CTCAGGCTCC CAGTTGGGAA -456

-455
CCAAGGAGGG GGAGGATGGG GGGGGGGGTG TGCGCCGACC CGGAAACGCC ATATAAGGAG -396

-395
CAGGAAGGAT CCCCCGCCGG AACAGACCTT ATTTGGGCAG CGCCTTATAT GGAGTGGCCC -336

FIGURE 4.1

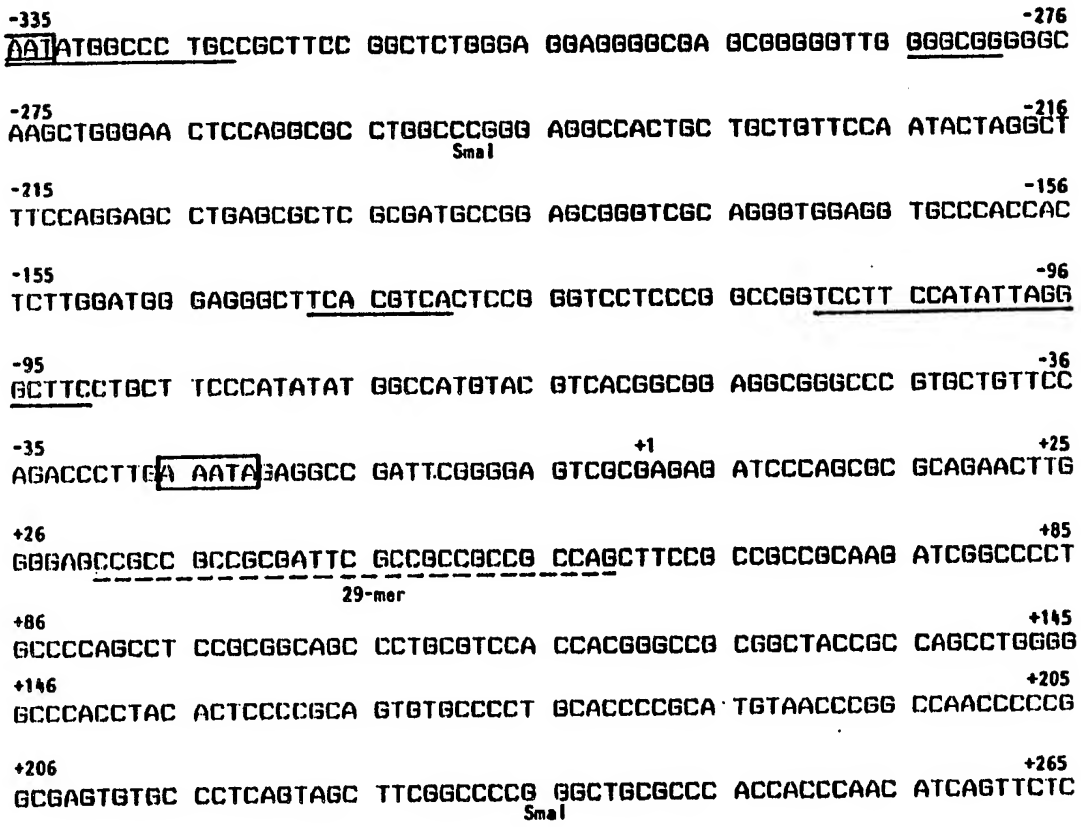

FIGURE 4.2

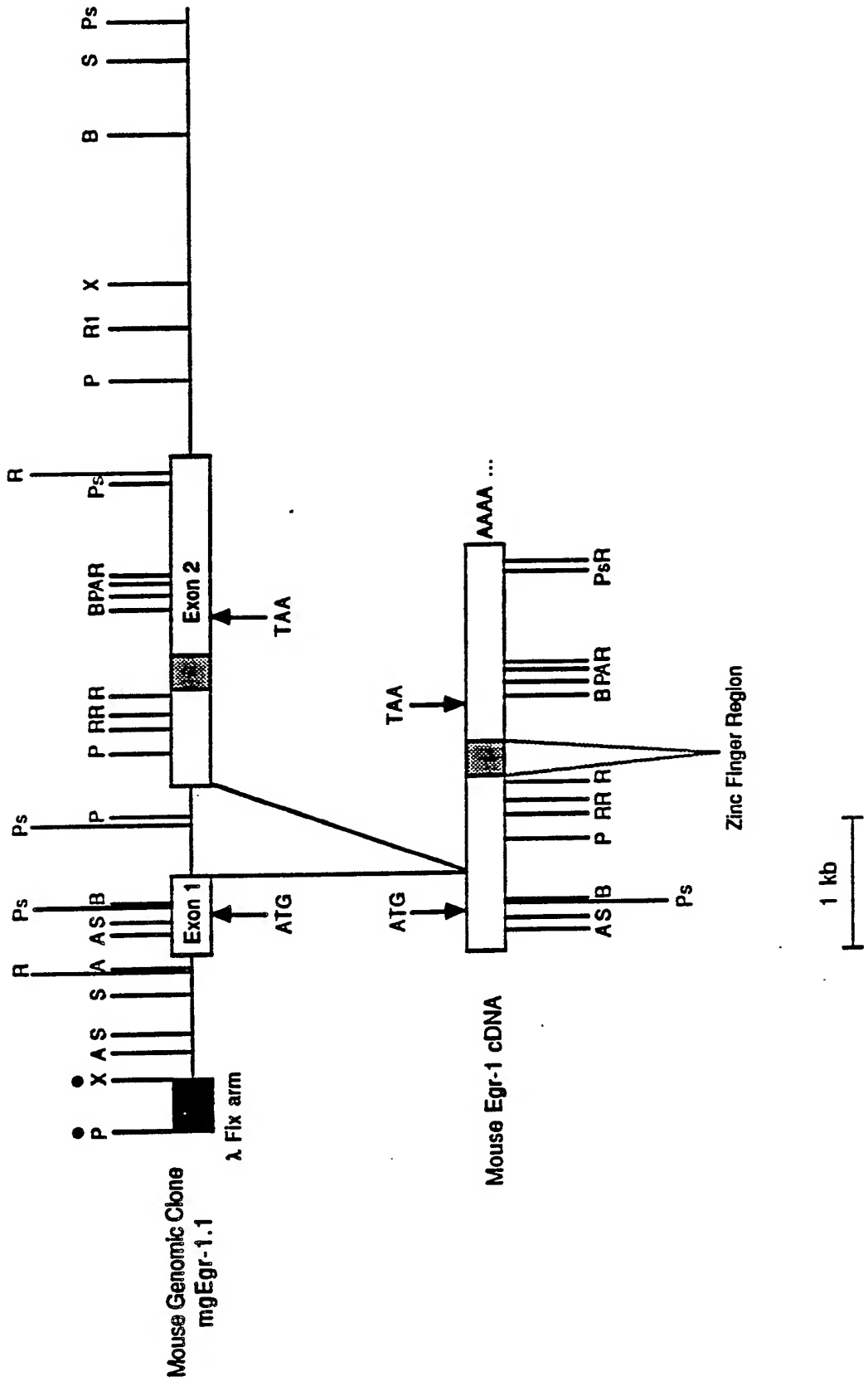
Mouse Genomic Clone
mgEgr-1.1
λ Fix arm
P
X
A S
S
A
R
Ps
B
A S
Exon 1
ATG
Ps
P
P RR R
BPAR
Exon 2
TAA
R
Ps
P
R1
X
B
S
Ps
Mouse Egr-1 cDNA
ATG
TAA
AAAA ...
AS
B
Ps
P RR R
BPAR
PsR
Zinc Finger Region
1 kb

FIGURE 5

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US89/01473

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) [6]

According to International Patent Classification (IPC) or to both National Classification and IPC

C07H 15/12, C12P 21/00, 19/34; C12N 15/00, 7/00, 1/20; C12Q 1/68; C07K 13/00

II. FIELDS SEARCHED

Minimum Documentation Searched [7]

Classification System	Classification Symbols
US	536/27; 435/68, 91, 172.1, 172.3, 320, 252.3, 6; 530/350, 387

Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched [8]

CAS database (1967-1989) keywords: finger/protein/motif early growth regulatory/gene/expression/sequence

III. DOCUMENTS CONSIDERED TO BE RELEVANT [9]

Category *	Citation of Document, [11] with indication, where appropriate, of the relevant passages [12]	Relevant to Claim No. [13]
X/Y	Chavrier, P. et al (January 1988) EMBO Journal, Volume 7: 29-35;" A gene encoding a protein with zinc fingers is activated during G_0/G_1, transition in cultured cells. See entire document.	1-5,8-10 / 6,7,11,12 14-16
X/Y	Chowdhury, K. et al (March 1988) Cell, Volume 48: 771-778; "A Multigene family encoding several finger structures is present and differentially active in mammalian genomes". See entire document.	1-5,8-10 / 6,7,11, 12,14-16
X/Y	Lau, L. et al (March 1987) PNAS, Volume 84: 1182-1186; "Expression of a set of growth-related immediate early genes in BALB/c 3T3 cells: coordinate regulation with c-fos or c-myc. See entire document	1-5,7-11 / 6,12, 14-16

* Special categories of cited documents: [10]

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

27 June 1989

Date of Mailing of this International Search Report

24 AUG 1989

International Searching Authority

ISA US

Signature of Authorized Officer

Anne Brown

Anne Brown

Form PCT/ISA/210 (second sheet) (Rev.11-87)

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

P,Y	Pannuti, A. et al (May 1988) Nucleic Acids Res., Volume 16: 4227-4237 "Isolation of CDNAs encoding finger proteins and measurment of the corresponding mRNA levels during myeloid terminal differentiation" See entire document.	1-12, 14-16
P,Y	Almendrol et al (May 1988) Mol Cell Biol, Vol. 8:2140-2148 "Complexity of the Early Genetic Response to Growth Factors in Mouse Fibroblasts". See entire document.	1-12, 14-16

V. ☐ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE [1]

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☐ Claim numbers . because they relate to subject matter [12] not required to be searched by this Authority, namely:

2. ☐ Claim numbers . . . , because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out [13], specifically:

3. ☐ Claim numbers_____, because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI ☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING [2]

This International Searching Authority found multiple inventions in this international application as follows:

See attached sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.

2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:

3. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers: 1-12, 14-16

4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

☐ The additional search fees were accompanied by applicant's protest.

☐ No protest accompanied the payment of additional search fees.

ATTACHMENT TO PCT/ISA/210
VI. OBSERVATION WHERE UNITY OF INVENTION IS LACKING

I. Claims 1-12, and 14-16 are drawn to EGR DNA sequence, vector or cell containing it, transformed host, method of using vector or cell and a polypeptide.

II. Claim 13 is drawn to cell-free method of preparing an early growth regulatory protein.

III. Claims 17-19 are drawn to native EGR protein.

IV. Claims 20-23 are drawn to synthetic peptide fragment antigenically related to and containing homology to native EGR protein.

V. Claims 24-29 are drawn to antibodies to EGR, anti-bodies to region EGR, method of using antibody.

VI. Claim 30 is drawn to method of detecting mRNA.

VII. Claim 31 is drawn to method of detecting DNA.

VIII. Claim 32 is drawn to method of diagnosis.

The claims do not embrace one single general inventive concept as defined in Rule 13.

Groups I, III, IV and V contain claims directed to distinct chemical entities. These are a gene and a protein coded for by that gene, a native EGR protein, a synthetic peptide fragment, antibodies which bind to EGR proteins. Groups II, VI, VII and VIII are directed to alternative uses for the nucleotide defined in Group I.